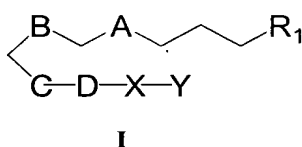


### Amendments to the Claims

This listing of claims will replace all prior versions and listings of claims in the application:

Claim 1 (currently amended). A composition for the treatment of dry eye and other disorders requiring the wetting of the eye comprising a pharmaceutically acceptable carrier and a pharmaceutically effective amount of one or more compounds of the following formula I:



wherein:

R<sup>1</sup> is CO<sub>2</sub>R, CONR<sup>2</sup>R<sup>3</sup>, CH<sub>2</sub>OR<sup>4</sup>, CH<sub>2</sub>NR<sup>5</sup>R<sup>6</sup>, CH<sub>2</sub>N<sub>3</sub>, CH<sub>2</sub>Hal, CH<sub>2</sub>NO<sub>2</sub>, CH<sub>2</sub>SR<sup>20</sup>, COSR<sup>21</sup>, or 2,3,4,5-tetrazol-1-yl, wherein:

R is H or CO<sub>2</sub>R forms a pharmaceutically acceptable salt or a pharmaceutically acceptable ester;

NR<sup>2</sup>R<sup>3</sup> and NR<sup>5</sup>R<sup>6</sup> are the same or different and comprise a free or functionally modified amino group, ~~e.g., R<sup>2</sup>, R<sup>3</sup>, R<sup>5</sup> and R<sup>6</sup> are the same or different and are H, alkyl, cycloalkyl, aralkyl, aryl, OH, or alkoxy,~~ with the proviso that at most only one of R<sup>2</sup> and R<sup>3</sup> ~~are~~ is OH or alkoxy and at most only one of R<sup>5</sup> and R<sup>6</sup> ~~are~~ is OH or alkoxy;

OR<sup>4</sup> comprises a free or functionally modified hydroxy group, ~~e.g., R<sup>4</sup> is H, acyl, alkyl, cycloalkyl, aralkyl, or aryl;~~

Hal is F, Cl, Br, or I;

SR<sup>20</sup> comprises a free or functionally modified thiol group; and

R<sup>21</sup> is H or COSR<sup>21</sup> forms a pharmaceutically acceptable salt or a pharmaceutically acceptable thioester;

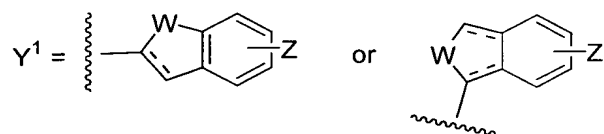
A, B and D are the same or different and are C<sub>1</sub>-C<sub>5</sub> alkyl, C<sub>2</sub>-C<sub>5</sub> alkenyl, C<sub>2</sub>-C<sub>5</sub> alkynyl, or a C<sub>3</sub>-C<sub>5</sub> allenyl group;

C is ~~an oxirane~~ or cyclopropane;

X is (CH<sub>2</sub>)<sub>m</sub> or (CH<sub>2</sub>)<sub>m</sub>O, wherein m is 1-6; and

Y is a phenyl ring optionally substituted with alkyl, halo, trihalomethyl, acyl, or a free or functionally modified hydroxy, amino, or thiol group; or

X-Y is (CH<sub>2</sub>)<sub>p</sub>Y<sup>1</sup>; wherein p is 0-6; and



wherein:

W is CH<sub>2</sub>, O, S(O)<sub>q</sub>, NR<sup>8</sup>, CH<sub>2</sub>CH<sub>2</sub>, CH=CH, CH<sub>2</sub>O, CH<sub>2</sub>S(O)<sub>q</sub>, CH=N, or CH<sub>2</sub>NR<sup>8</sup>; wherein q is 0-2, and R<sup>8</sup> is H, alkyl, or acyl;

Z is H, alkyl, acyl, halo, trihalomethyl, or a free or functionally modified amino, thiol, or hydroxy group; and

==== is a single or double bond;

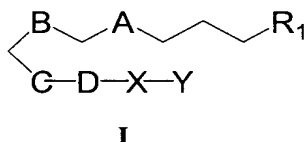
or X-Y is cyclohexyl or *n*-C<sub>5</sub>H<sub>11</sub>.

Claim 2 (cancelled).

Claim 3 (cancelled).

Claim 4 (original). The composition of Claim 1, wherein the composition is a topical ophthalmic formulation.

Claim 5 (currently amended). A method for the treatment of dry eye and other disorders requiring the wetting of the eye which comprises administering to a mammal a composition comprising a pharmaceutically acceptable carrier and a pharmaceutically effective amount of one or more compounds of the following formula I:



wherein:

R<sup>1</sup> is CO<sub>2</sub>R, CONR<sup>2</sup>R<sup>3</sup>, CH<sub>2</sub>OR<sup>4</sup>, CH<sub>2</sub>NR<sup>5</sup>R<sup>6</sup>, CH<sub>2</sub>N<sub>3</sub>, CH<sub>2</sub>Hal, CH<sub>2</sub>NO<sub>2</sub>, CH<sub>2</sub>SR<sup>20</sup>, COSR<sup>21</sup>, or 2,3,4,5-tetrazol-1-yl, wherein:

R is H or CO<sub>2</sub>R forms a pharmaceutically acceptable salt or a pharmaceutically acceptable ester;

NR<sup>2</sup>R<sup>3</sup> and NR<sup>5</sup>R<sup>6</sup> are the same or different and comprise a free or functionally modified amino group, e.g., R<sup>2</sup>, R<sup>3</sup>, R<sup>5</sup> and R<sup>6</sup> are the same or different and are H, alkyl, cycloalkyl, aralkyl, aryl, OH, or alkoxy, with the proviso that at most only one of R<sup>2</sup> and R<sup>3</sup> are is OH or alkoxy and at most only one of R<sup>5</sup> and R<sup>6</sup> are is OH or alkoxy;

OR<sup>4</sup> comprises a free or functionally modified hydroxy group, e.g., R<sup>4</sup> is H, acyl; alkyl, cycloalkyl, aralkyl, or aryl;

Hal is F, Cl, Br, or I;

SR<sup>20</sup> comprises a free or functionally modified thiol group; and

$R^{21}$  is H or  $\text{COSR}^{21}$  forms a pharmaceutically acceptable salt or a pharmaceutically acceptable thioester;

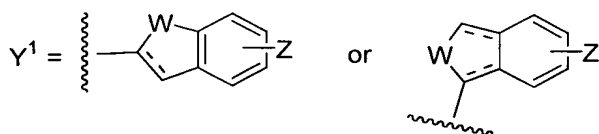
A, B and D are the same or different and are  $\text{C}_1\text{-C}_5$  alkyl,  $\text{C}_2\text{-C}_5$  alkenyl,  $\text{C}_2\text{-C}_5$  alkynyl, or a  $\text{C}_3\text{-C}_5$  allenyl group;

C is an oxirane or cyclopropane;

X is  $(\text{CH}_2)_m$  or  $(\text{CH}_2)_m\text{O}$ , wherein m is 1-6; and

Y is a phenyl ring optionally substituted with alkyl, halo, trihalomethyl, acyl, or a free or functionally modified hydroxy, amino, or thiol group; or

X-Y is  $(\text{CH}_2)_p\text{Y}^1$ ; wherein p is 0-6; and



wherein:

W is  $\text{CH}_2$ , O,  $\text{S(O)}_q$ ,  $\text{NR}^8$ ,  $\text{CH}_2\text{CH}_2$ ,  $\text{CH=CH}$ ,  $\text{CH}_2\text{O}$ ,  $\text{CH}_2\text{S(O)}_q$ ,  $\text{CH=N}$ , or  $\text{CH}_2\text{NR}^8$ ; wherein q is 0-2, and  $\text{R}^8$  is H, alkyl, or acyl;

Z is H, alkyl, acyl, halo, trihalomethyl, or a free or functionally modified amino, thiol, or hydroxy group; and

---- is a single or double bond;

or X-Y is cyclohexyl or  $n\text{-C}_5\text{H}_{11}$ .

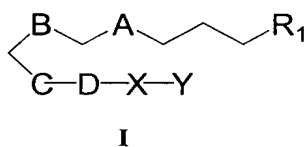
Claim 6 (cancelled).

Claim 7 (cancelled).

Claim 8 (original). The method of Claim 5, wherein the composition is a topical ophthalmic formulation.

Claim 9 (original). The method of Claim 5 wherein the dry eye and other disorders requiring the wetting of the eye is symptoms of dry eye associated with refractive surgery.

Claim 10 (currently amended). A compound of the following formula I:



wherein:

R<sup>1</sup> is CO<sub>2</sub>R, CONR<sup>2</sup>R<sup>3</sup>, CH<sub>2</sub>OR<sup>4</sup>, CH<sub>2</sub>NR<sup>5</sup>R<sup>6</sup>, CH<sub>2</sub>N<sub>3</sub>, CH<sub>2</sub>Hal, CH<sub>2</sub>NO<sub>2</sub>, CH<sub>2</sub>SR<sup>20</sup>, COSR<sup>21</sup>, or 2,3,4,5-tetrazol-1-yl, wherein:

R is H or CO<sub>2</sub>R forms a pharmaceutically acceptable salt or a pharmaceutically acceptable ester;

NR<sup>2</sup>R<sup>3</sup> and NR<sup>5</sup>R<sup>6</sup> are the same or different and comprise a free or functionally modified amino group, ~~e.g., R<sup>2</sup>, R<sup>3</sup>, R<sup>5</sup> and R<sup>6</sup> are the same or different and are H,~~ alkyl, cycloalkyl, aralkyl, aryl, OH, or alkoxy, with the proviso that at most only one of R<sup>2</sup> and R<sup>3</sup> are is OH or alkoxy and at most only one of R<sup>5</sup> and R<sup>6</sup> are is OH or alkoxy;

OR<sup>4</sup> comprises a free or functionally modified hydroxy group, ~~e.g., R<sup>4</sup> is H, acyl,~~ alkyl, cycloalkyl, aralkyl, or aryl;

Hal is F, Cl, Br, or I;

SR<sup>20</sup> comprises a free or functionally modified thiol group; and

$R^{21}$  is H or  $\text{COSR}^{21}$  forms a pharmaceutically acceptable salt or a pharmaceutically acceptable thioester;

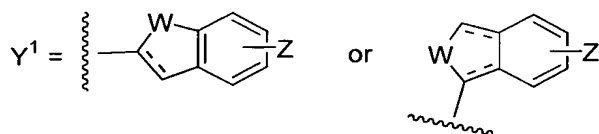
A, B and D are the same or different and are  $\text{C}_1\text{-C}_5$  alkyl,  $\text{C}_2\text{-C}_5$  alkenyl,  $\text{C}_2\text{-C}_5$  alkynyl, or a  $\text{C}_3\text{-C}_5$  allenyl group;

C is an oxirane or cyclopropane;

X is  $(\text{CH}_2)_m$  or  $(\text{CH}_2)_m\text{O}$ , wherein m is 1-6; and

Y is a phenyl ring optionally substituted with alkyl, halo, trihalomethyl, acyl, or a free or functionally modified hydroxy, amino, or thiol group; or

X-Y is  $(\text{CH}_2)_p\text{Y}^1$ ; wherein p is 0-6; and



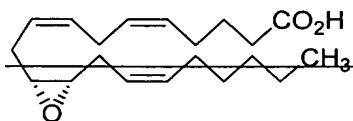
wherein:

W is  $\text{CH}_2$ , O,  $\text{S(O)}_q$ ,  $\text{NR}^8$ ,  $\text{CH}_2\text{CH}_2$ ,  $\text{CH=CH}$ ,  $\text{CH}_2\text{O}$ ,  $\text{CH}_2\text{S(O)}_q$ ,  $\text{CH=N}$ , or  $\text{CH}_2\text{NR}^8$ ; wherein q is 0-2, and  $\text{R}^8$  is H, alkyl, or acyl;

Z is H, alkyl, acyl, halo, trihalomethyl, or a free or functionally modified amino, thiol, or hydroxy group; and

---- is a single or double bond;

or X-Y is cyclohexyl or  $n\text{-C}_5\text{H}_{11}$ , provided that the following compound is excluded:



Claim 11 (cancelled).

Claim 12 (cancelled).